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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/748,765	12/29/2003	Illana Gozes	019856-000210US	8714
	7590 10/28/200 AND TOWNSEND AN	EXAMINER		
TWO EMBAR	CADERO CENTER	WOODWARD, CHERIE MICHELLE		
EIGHTH FLOO SAN FRANCIS	SCO, CA 94111-3834	ART UNIT	PAPER NUMBER	
			1647	
			MAIL DATE	DELIVERY MODE
			10/28/2008	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

## Advisory Action Before the Filing of an Appeal Brief

Application No.	Applicant(s)	
10/748,765	GOZES ET AL.	
Examiner	Art Unit	
CHERIE M. WOODWARD	1647	

	CHERIE M. WOODWARD	1647	
The MAILING DATE of this communication appe	ars on the cover sheet with the c	correspondence add	ress
THE REPLY FILED 29 September 2008 FAILS TO PLACE THI	S APPLICATION IN CONDITION F	OR ALLOWANCE.	
1.  The reply was filed after a final rejection, but prior to or on application, applicant must timely file one of the following application in condition for allowance; (2) a Notice of Appetor Continued Examination (RCE) in compliance with 37 C periods:	replies: (1) an amendment, affidavit eal (with appeal fee) in compliance v	t, or other evidence, w with 37 CFR 41.31; or	hich places the (3) a Request
a) $\boxtimes$ The period for reply expires $\underline{5}$ months from the mailing date	of the final rejection.		
b) The period for reply expires on: (1) the mailing date of this A no event, however, will the statutory period for reply expire to Examiner Note: If box 1 is checked, check either box (a) or (MONTHS OF THE FINAL REJECTION. See MPEP 706.07(1)	ater than SIX MONTHS from the mailing b). ONLY CHECK BOX (b) WHEN THE f).	g date of the final rejection FIRST REPLY WAS FIL	n. .ED WITHIN TWO
Extensions of time may be obtained under 37 CFR 1.136(a). The date have been filed is the date for purposes of determining the period of extunder 37 CFR 1.17(a) is calculated from: (1) the expiration date of the set forth in (b) above, if checked. Any reply received by the Office later may reduce any earned patent term adjustment. See 37 CFR 1.704(b). NOTICE OF APPEAL	ension and the corresponding amount on hortened statutory period for reply origing than three months after the mailing date	of the fee. The appropria nally set in the final Offic	ite extension fee e action; or (2) as
2. The Notice of Appeal was filed on A brief in comp filing the Notice of Appeal (37 CFR 41.37(a)), or any exter Notice of Appeal has been filed, any reply must be filed with AMENDMENTS	nsion thereof (37 CFR 41.37(e)), to	avoid dismissal of the	
3. X The proposed amendment(s) filed after a final rejection, b	out prior to the data of filing a brief	will not be entered be	031160
(a) ☐ They raise new issues that would require further cor (b) ☐ They raise the issue of new matter (see NOTE belogical of the policy).  (c) ☐ They are not deemed to place the application in bet	nsideration and/or search (see NOT w);	E below);	
appeal; and/or			
(d) They present additional claims without canceling a c		ected claims.	
NOTE: <u>See Continuation Sheet</u> . (See 37 CFR 1.1 4. The amendments are not in compliance with 37 CFR 1.12	. , ,	mpliant Amondment (F	OTOL 324)
<ul> <li>5. Applicant's reply has overcome the following rejection(s):</li> </ul>		inpliant Amendment (r	10L-324).
<ol> <li>Newly proposed or amended claim(s) would be all non-allowable claim(s).</li> </ol>		imely filed amendmer	t canceling the
7.  For purposes of appeal, the proposed amendment(s): a) how the new or amended claims would be rejected is proved the status of the claim(s) is (or will be) as follows: Claim(s) allowed:		l be entered and an ex	xplanation of
Claim(s) allowed: Claim(s) objected to:			
Claim(s) rejected: <u>1,10-15,17-22 and 26-28</u> . Claim(s) withdrawn from consideration:			
AFFIDAVIT OR OTHER EVIDENCE  No. The efficient or other evidence filed effects final action, but	t before or on the date of filing a Ne	tice of Annacl will not	he entered
8.  The affidavit or other evidence filed after a final action, bubecause applicant failed to provide a showing of good and was not earlier presented. See 37 CFR 1.116(e).			
<ol> <li>The affidavit or other evidence filed after the date of filing entered because the affidavit or other evidence failed to o showing a good and sufficient reasons why it is necessary</li> </ol>	vercome <u>all</u> rejections under appea and was not earlier presented.  Se	ıl and/or appellant fails ee 37 CFR 41.33(d)(1)	s to provide a
10. The affidavit or other evidence is entered. An explanation	n of the status of the claims after er	ntry is below or attache	ed.
REQUEST FOR RECONSIDERATION/OTHER  11. The request for reconsideration has been considered but .	t does NOT place the application in	condition for allowand	ce because:
12. Note the attached Information <i>Disclosure Statement</i> (s). (13. Other:	PTO/SB/08) Paper No(s)		
/Gary B. Nickol / Supervisory Patent Examiner, Art Unit 1646			

Continuation of 3. NOTE: Applicant's arguments directed to the written description rejection under 35 USC 112, first paragraph have been considered, but they are not persuasive. Applicant's case law arguments are spurious when the examiner has shown that the genus of ADNF III polypeptides and the genus of ADNF I polypeptides are highly variable in structure (see NCBI references recited in the Office Action of 6 July 2006). In order to comply with the written description requirement, the structure which is asserted to make up the polypeptide must be clearly and positively specified. The structure must be organized and correlated in such a manner as to present a complete operative embodiment which is adequately described in the specification. The instant disclosure fails to provide an adequate description of a sufficient number of variant ADNF III and ADNF I polypeptides that function to treat MS. The general knowledge and level of those of ordinary skill does not supplement the omitted description because specific, not general, descriptions are needed.

Regarding Applicant's arguments directed to the rejections under 35 USC 103a, have been considered, but they are not persuasive. Applicant's reliance on In re Riickaert, 28 USPQ2d 1955 (Fed. Cir. 1993) is misplaced. In Riickaert, the Court relied on In re Oelright, 666 F.2d 578, 581-2, 212 USPQ 323, 326 (CCPA 1981), which held that "[t]he mere fact that a certain thing may result from a given set of circumstances is not sufficient [to establish inherency." Additionally, the Rijckaert Court relied on In re Spormann, 363 F.2d 444, 448, 150 USPQ 449, 452 (CCPA 1966), which held "[t]hat which may be inherent is not necessarily known. Obviousness cannot be predicated on what is unknown." The Court held that such a retrospective view of inherency is not a substitute for some teaching or suggestion supporting an obviousness rejection, citing In re Newell, 891 F.2d 899, 901, 13 USPQ2d 1248, 1250 (Fed.Cir. 1989). The facts, statements, and holdings of the Rijckaert Court factually distinct from the facts of record in the instant case. The examiner has expressly stated of record that the '740 patent and WO 98/35042 teach the administration ADNF peptides as therapeutics to treat neurodegenerative disorders, including Guillan-Barre syndrome, and Brenneman et al., teach the use of ADNF polypeptides to treat conditions related to increased neuronal cell death. As stated in the Office Actions of record, at the time of the invention, there was a recognized problem or need in the art to treat multiple sclerosis. There were a finite number of identified, predictable potential solutions to treat related neurological and autoimmune disorders using an ADNF polypeptide or active core sequence thereof. One of ordinary skill in the art could have pursued the known potential solutions with a reasonable expectation of success because the '740 patent and WO 98/35042 teach the use of ADNF polypeptides and active core sequences thereof for neurological and autoimmune disorders and Brenneman et al., teach the administration of ADNF polypeptides to treat conditions related to increased neuronal cell death. A person of ordinary skill in the art at the time the invention was made would have reasonably known that the ADNF polypeptides and active core sites thereof would have useful in the treatment of neurological disorders, including autoimmune neurological disorders, and would also be useful in the treatment of multiple sclerosis. Moreover, WO 98/35042 teaches that those of skill in the art will appreciate that the list of neurodegenerative disorders is not exhaustive and that ADNF III polypeptides can be used to treat other neurological disorders (page 8, lines 16-18). The teachings of the prior art, as cited by the examiner, are sufficient to permit a person of ordinary skill in the art to recognize the inherent functions of the ADNF polypeptides. Moreover, the teachings of the prior art provide the rationale and motivations to choose from a finite number of identified, predictable solutions, with a reasonable expectation of success (see MPEP 2141(III) Rationale E, also recited as Examination Guidelines for Determining Obviousness under 35 USC 103 in view of the Supreme Court Decision in KSR International Co. v. Teleflex, Inc., as set forth of record).

The submission of the self-serving declaration of a co-inventor under 37 CFR 1.132, where the declarant attempts to demonstrate that the prior art references do not provide motivation for their combination, is not persuasive. The rationale and motivation in the prior art references speak for themselves, as discussed in detail in the rejections of record. Further, Applicant has provided this declaration for the first time in an After-Final submission. There is no showing of good and sufficient reasons why the affidavit was not earlier presented (see 37 CFR 1.116(e)). The declaration is not entered of record.

Regarding Applicant's arguments as to the Double Patenting Rejection, Applicant's arguments have been considered, but they are not persuasive. Applicant argues that a two-way test for obviousness is required over the 11/388,634 ('634) application (now allowed, issued fee paid 10/3/2008). The '634 application has a common inventor and a common assignee with the instant application and claims 1 and 23 of the '634 application are not patentably distinct from instant claims 1, 11, 14, 17, 20, and 21, for the reasons of record. The reference is a pending, but allowed patent application. As such, a one-way test applies. It is noted that Applicant did not pay the issue fee in the '634 case until 10/3/2008, several days after filing the instant After-Final request and that the '634 application has not yet in fact issued as a patent. Arguably, even if the '634 application had previously issued as a patent, a two-way test would still not apply because although the instant application is the earlier filed of the two, Applicant has presented no evidence, showing, or petition decision regarding any administrative delay on the part of the Office causing delay in prosecution of the application. Moroever, Applicant has not made a showing or prodvided evidence as to why the conflicting claims could not have been filed in a single application. Both administrative delay on the part of the Office and the showing that the conflicting claims could not have been filed in a single application must be shown in order to apply the two-way test (see MPEP 804).

It is noted that Applicant's arguments (page 15 of 17, first paragraph) states that the pending application claims priority to a provisional application and a PCT application, the national stage of which was filed on 29 December 2003. This is factually incorrect. The instant application only claims benefit under 35 USC 119(e) to US provisional application 60/437650, filed 1/2/2003.

The rejections of record are maintained for the reasons of record and the reasons set forth herein. /CMW/ AU 1647